Listing of Claims:

 (Currently Amended) Benzimidazole derivatives A compound or compounds of formula I

$$(R^8)_p$$
 N
 R^7
 X
 Ar^2
 $(R^{10})_r$
 R^6
 $(R^9)_q$

wherein

 R^6 , R^7 are independently from one another H, A or SO_2A ,

A is independently selected from the group consisting of alkyl, alkenyl, cycloalkyl, alkylenecycloalkyl, alkoxy and alkoxyalkyl,

Ar² is selected independently from one another from aromatic hydrocarbons containing 6 to 14 carbon atoms and ethylenical unsaturated or aromatic heterocyclic residues containing 3 to 10 carbon atoms and one or two heteroatoms, independently selected from the group consisting of N, O and S,

 R^8 , R^9 and R^{10} are independently selected from a <u>the</u> group consisting of H, A, cycloalkyl comprising 3 to 7 carbon atoms, Hal, CH₂Hal, CH(Hal)₂, C(Hal)₃, NO₂, (CH₂)_nCN, (CH₂)_nNR¹¹R¹², (CH₂)_nOR¹¹, (CH₂)_nO(CH₂)_kNR¹¹R¹², (CH₂)_nCOOR¹², (CH₂)_nCONR¹¹R¹², (CH₂)_nNR¹¹CONR¹¹R¹², (CH₂)_nNR¹¹CONR¹¹R¹², (CH₂)_nNR¹¹SO₂A, (CH₂)_nSO₂NR¹¹R¹², (CH₂)_nS(O)_uR¹³,

(CH₂)_nOC(O)R¹³, (CH₂)_nCOR¹³, (CH₂)_nSR¹¹, CH=N-OA, CH₂CH=N-OA, (CH₂)_nNHOA, (CH₂)_nCH=N-R¹¹, (CH₂)_nOC(O)NR¹¹R¹², (CH₂)_nNR¹¹COOR¹², (CH₂)_nN(R¹¹)CH₂CH₂OR¹³, (CH₂)_nN(R¹¹)CH₂CH₂OCF₃, (CH₂)_nN(R¹¹)C(R¹³)HCOOR¹², C(R¹³)HCOR¹², (CH₂)_nN(R¹¹)CH₂CH₂N(R¹²)CH₂COOR¹², (CH₂)_nN(R¹¹)CH₂CH₂NR¹¹R¹², CH=CHCOOR¹¹, CH=CHCH₂NR¹¹R¹², CH=CHCH₂NR¹¹R¹², CH=CHCH₂OR¹³, (CH₂)_nN(COOR¹¹)COOR¹², (CH₂)_nN(CONH₂)COOR¹¹, (CH₂)_nN(CONH₂)CONH₂, (CH₂)_nN(CH₂COOR¹¹)COOR¹², (CH₂)_nN(CH₂CONH₂)COOR¹¹, (CH₂)_nN(CH₂CONH₂)CONH₂, (CH₂)_nCHR¹³COR¹¹, (CH₂)_nCHR¹³COOR¹¹, (CH₂)_nCHR¹³COR¹¹, (CH₂)_nCHR¹³COOR¹¹,

- R^{11} , R^{12} are independently selected from a <u>the</u> group consisting of H, A, $(CH_2)_mAr^3$ and $(CH_2)_mHet$, or in $NR^{11}R^{12}$,
- R¹¹ and R¹² form, together with the N-atom they are bound to, a 5-, 6- or 7-membered heterocyclus heterocycle which optionally contains 1 or 2 additional heteroatoms, selected from the group consisting of N, O and S,
- R^{13} , R^{14} are independently selected from a <u>the</u> group consisting of H, Hal, A, $(CH_2)_mAr^4$ and $(CH_2)_mHet$,
- Ar³, Ar⁴ are independently from one another aromatic hydrocarbon residues comprising 5 to 12 and preferably 5 to 10 carbon atoms which are optionally substituted by one or more substituents, selected from a the group consisting of A, Hal, NO₂, CN, OR¹⁵, NR¹⁵R¹⁶, COOR¹⁵, CONR¹⁵R¹⁶, NR¹⁵COR¹⁶, NR¹⁵CONR¹⁵R¹⁶,

NR¹⁶SO₂A, COR¹⁵, SO₂R¹⁵R¹⁶, S(O)_uA and OOCR¹⁵,

Het is a saturated, unsaturated or aromatic heterocyclic residue which is optionally substituted by one ore or more substituents, selected from a the group consisting of A, Hal, NO₂, CN, OR¹⁵, NR¹⁵R¹⁶, COOR¹⁵, CONR¹⁵R¹⁶, NR¹⁵COR¹⁶, NR¹⁵CONR¹⁵R¹⁶, NR¹⁶SO₂A, COR¹⁵, SO₂R¹⁵R¹⁶, S(O)_uA and OOCR¹⁵,

R¹⁵, R¹⁶ are independently selected from a <u>the</u> group consisting of H, A, and (CH₂)_mAr⁶, wherein

Ar⁶ is a 5- or 6-membered aromatic hydrocarbon which is optionally substituted by one or more substituents selected from a <u>the</u> group consisting of methyl, ethyl, propyl, 2-propyl, tert.-butyl, Hal, CN, OH, NH₂ and CF₃,

k, m and n are independently of one another 0, 1, 2, 3, 4, or 5,

- X represents a bond or is $(CR^{11}R^{12})_h$, or $(CHR^{11})_h$ -Q- $(CHR^{12})_i$, wherein
- Q is selected from a the group consisting of O, S, N-R¹⁵, (CHal₂)_j, (O-CHR¹⁸)_j, (CHR¹⁸-O)_j, CR¹⁸=CR¹⁹, (O-CHR¹⁸CHR¹⁹)_j, (CHR¹⁸CHR¹⁹-O)_j, C=O, C=S, C=NR¹⁵, CH(OR¹⁵), C(OR¹⁵)(OR²⁰), C(=O)O, OC(=O), OC(=O)O, C(=O)N(R¹⁵), N(R¹⁵)C(=O), OC(=O)N(R¹⁵), N(R¹⁵)C(=O)O, CH=N-O, CH=N-NR¹⁵, S=O, SO₂, SO₂NR¹⁵ and NR¹⁵SO₂, wherein
- R^{18} , R^{19} , R^{20} are independently selected from the meanings given for R^8 , R^9 and R^{10} , preferably independently selected from the group

consiting of H, A, Hal, CH ₂ Hal, CH(Hal) ₂ , C(Hal) ₃ , NO ₂ ,
$(CH_2)_nCN, (CH_2)_nOR^{11}, (CH_2)_nNR^{11}R^{12},$
$(CH_2)_nO(CH_2)_kNR^{11}R^{12}, (CH_2)_nCOOR^{13}, (CH_2)_nCONR^{11}R^{12},$
$(CH_2)_nNR^{11}COR^{13}, (CH_2)_nNR^{11}CONR^{11}R^{12}, (CH_2)_nNR^{11}SO_2A,$
$(CH_2)_nSO_2NR^{11}R^{12}, (CH_2)_nS(O)_uR^{13}, (CH_2)_nCOR^{13}, (CH_2)_nSR^{11},$
(CH ₂) _n NHOA and (CH ₂) _n NR ¹¹ COOR ¹³ ,
·

- h, i are independently from each other 0, 1, 2, 3, 4, 5, or 6, and
- j is 1, 2, 3, 4, 5, or 6,
- Y is selected from the group consisting of O, S, NR^{21} , $C(R^{22})$ - NO_2 , $C(R^{22})$ -CN and $C(CN)_2$, wherein
- R^{21} is independently selected from the meanings given for R^{13} , R^{14} and
- R^{22} is independently selected from the meanings given for R^{11} , R^{12} ,
- p, r are independently from one another 0, 1, 2, 3, 4 or 5,
- q is 0, 1, 2, 3 or 4, preferably 0, 1 or 2,
- u is 0, 1, 2 or 3, preferably 0, 1 or 2,

and

Hal is independently selected from a <u>the</u> group consisting of F, Cl,

Br and I; <u>or</u>

and the physiologically acceptable derivatives, salts, and solvates thereof or mixtures thereof in all ratios.

- 2. (Currently Amended) Benzimidazole derivative A compound or compounds according to claim 1, wherein
 - Ar² is selected from aromatic hydrocarbons containing 6 to 10 and especially 6 carbon atoms and ethylenical unsaturated or aromatic heterocyclic residues containing 3 to 8 and especially 4 to 6 carbon atoms and one or two heteroatoms, independently selected from the group consisting of N, O and S and especially selected from N and O,
 - R⁸, R⁹ and R¹⁰ are independently selected from a the group consisting of H, A, cycloalkyl 3 to 7 carbon atoms, Hal, CH₂Hal, CH(Hal)₂, C(Hal)₃, NO₂, (CH₂)_nCN, (CH₂)_nOR¹¹, (CH₂)_nNR¹¹R¹², (CH₂)_nCONR¹¹R¹², (CH₂)_nO(CH₂)_kNR¹¹R¹², (CH₂)_nCOOR¹³, (CH₂)_nNR¹¹CONR¹¹R¹², (CH₂)_nNR¹¹SO₂A, (CH₂)_nSO₂NR¹¹R¹², (CH₂)_nNR¹¹CONR¹¹R¹², (CH₂)_nOC(O)R¹³, (CH₂)_nCOR¹³, (CH₂)_nS(O)_uR¹³, (CH₂)_nNHOA, (CH₂)_nNR¹¹COOR¹³, (CH₂)_nN(R¹¹)CH₂CH₂OCF₃, (CH₂)_nN(R¹¹)CH₂CH₂OR¹³, (CH₂)_nN(R¹¹)CH₂CH₂OCF₃, (CH₂)_nN(R¹¹)C(R¹³)HCOOR⁸, (CH₂)_nN(CONH₂)COOR¹³, (CH₂)_nN(COOR¹³)COOR¹⁴, (CH₂)_nN(CONH₂)COOR¹³, (CH₂)_nN(CH₂COOR¹³)COOR¹⁴, (CH₂)_nN(CH₂COOR¹³)COOR¹⁴, (CH₂)_nN(CH₂CONH₂)COOR¹⁴, (CH₂)_nN(CH₂CONH₂)CONH₂, (CH₂)_nCHR¹³COR¹⁴, (CH₂)_nCHR¹³COR¹⁴, (CH₂)_nCHR¹³COOR¹⁴ and (CH₂)_nCHR¹³COR¹⁴,
 - represents a bond or is $(CR^{11}R^{12})_h$, or $(CHR^{11})_h$ -Q- $(CHR^{12})_i$,

wherein

Q is selected from a the group consisting of O, S, N-R¹⁵, (CHal₂)_j, (O-CHR¹⁸)_j, (CHR¹⁸-O)_j, CR¹⁸=CR¹⁹, (O-CHR¹⁸CHR¹⁹)_j, (CHR¹⁸CHR¹⁹-O)_j, C=O, C=NR¹⁵, CH(OR¹⁵), C(OR¹⁵)(OR²⁰), C(=O)N(R¹⁵), N(R¹⁵)C(=O), CH=N-NR¹⁵, S=O, SO₂, SO₂NR¹⁵ and NR¹⁵SO₂, wherein

h, i are independently from each other 0, 1, 2, 3, 4, 5 or 6, preferably 0, 1, 2 or 3 and

j is 1, 2, 3, 4, 5 or 6, preferably 1, 2, 3 or 4,

p is 1, 2, 3 or 4, preferably 1, 2 or 3, and

r is 0, 1, 2, or 3, preferably 0, 1 or 2;

and the physiologically acceptable derivatives, salts and solvates thereof.

3. (Currently Amended) Benzimidazole derivative The compound or compounds according to claim 1, selected from the group consisting of the compounds of the formulae Ia, Ib, Ic and Id,

$$(R^8)_p$$
 N
 R^7
 R^{10}
 R^{10}

$$(R^8)_p$$
 N
 N
 R^7
 R^{10}
 R^{10}
 R^{10}

$$(R^8)_p \xrightarrow{\stackrel{\textstyle H}{\textstyle N}} \stackrel{\textstyle R^7}{\textstyle N} \xrightarrow{\textstyle R^{10}}$$

$$(R^9)_q$$
Ic

$$(R^8)_p \xrightarrow{\stackrel{\textstyle H}{\stackrel{\textstyle N}{\stackrel{\textstyle N}{\stackrel {\stackrel \textstyle N}{\stackrel \textstyle N}{\stackrel \textstyle N}{\stackrel \textstyle N}}}}}} Id}$$

wherein

R⁷, R⁸, p, X, Y, R⁹ and q are as defined in claim 1, and R¹⁰ is H or as defined in claim 1;

and the physiologically acceptable derivatives, salts, and solvates thereof and mixtures thereof in all ratios.

4. (Currently Amended) Benzimidazole derivative The compound or compounds according to claim 3, additionally comprising one or two substituents selected from the group consisting of O(CH₂)_nNR¹¹R¹², NR¹¹(CH₂)_nNR¹¹R¹², O(CH₂)_nOR¹² and NR¹¹(CH₂)_nOR¹²,

wherein

- R^{11} , R^{12} are independently selected from a <u>the</u> group consisting of H, A, $(CH_2)_mAr^3$ and $(CH_2)_mHet$, or in $NR^{11}R^{12}$,
- R¹¹ and R¹² form, together with the N-atom they are bound to, a 5-, 6- or
 7-membered heterocyclus heterocycle which said heterocycle
 optionally contains 1 or 2 additional heteroatoms, selected from
 the group consisting of N, O and S, and
- n is 1, 2, 3, 4, 5 or 6.
- 5. (Currently Amended) Benzimidazole derivative The compound or compounds according to claim 1, selected from the group consisting of the compounds (1) to (78) of table 1; and the physiologically acceptable derivatives, salts, and solvates thereof and mixtures thereof in all ratios.
- 6. (Currently Amended) Benzimidazole derivative The compound or compounds according to claim 1 as a medicament.
- 7. (Currently Amended) Benzimidazole derivative The compound or compounds according to claim 1 as a kinase inhibitor.
- 8. (Currently Amended) Benzimidazole derivative The compound or compounds according to claim 7, characterized in that the kinases are selected from the group consisting of raf-kinases and VEGFR kinases.
- 9. (Currently Amended) A pharmaceutical Pharmaceutical composition, characterized in that it contains comprising one or more of the compound or compounds according to claim 1 as a pharmaceutical composition.

- 10. (Currently Amended) The pharmaceutical Pharmaceutical composition according to claim 9, characterised characterized in that it contains one or more additional compounds, selected from the group consisting of physiologically acceptable excipients, auxiliaries, adjuvants, carriers and pharmaceutical active ingredients.
- 11. (Currently Amended) Process A process for the manufacture of a pharmaceutical composition, characterised comprising in that one or more of the compound or compounds according to claim 1 and one or more compound or compounds, selected from the group consisting of carriers, excipients, auxiliaries and pharmaceutical active ingredients other than the compound or compounds according to claim 1, is processed by mechanical means into a pharmaceutical composition that is suitable as dosage_form for application and/or administration to a patient.
- 12. (Currently Amended) Use A method of comprising administering to a patient a the compound or compounds according to claim 1 as a pharmaceutical composition.
- 13. (Currently Amended) Use A method of comprising administering to a patient a the compound or compounds according to claim 1 in the treatment and/or prophylaxis of a disorder or disorders.
- 14. (Cancelled)
- 15. (Currently Amended) Use according to The method of claim 13, characterised characterized in that the disorder or disorders are caused, mediated and/or propagated by kinases selected from the group consisting of raf-kinases and VEGFR kinases.

- 16. (Currently Amended) Use according to The method of claim 13, characterised characterized in that the disorder or disorders are selected from the group consisting of hyperproliferative and nonhyperproliferative disorders.
- 17. (Currently Amended) Use according to The method of claim 13, characterised characterized in that the disorder or disorders is cancer.
- 18. (Currently Amended) Use according to The method of claim 13, characterised characterized in that the disorder or disorders is noncancerous.
- 19. (Currently Amended) Use according to The method of claim 18, characterised characterized in that the noncancerous disorder or disorders are selected from the group consisting of infection, psoriasis, arthritis, inflammation, endometriosis, scarring, begnin benign prostatic hyperplasia, immunological diseases, autoimmune diseases and immunodeficiency diseases.
- 20. (Currently Amended) Use according to The method of claim 17, characterised characterized in that the disorder or disorders are selected from the group consisting of brain cancer, lung cancer, squamous cell cancer, bladder cancer, gastric cancer, pancreatic cancer, hepatic cancer, renal cancer, colorectal cancer, breast cancer, head cancer, neck cancer, oesophageal cancer, gynaecological cancer, thyroid cancer, lymphoma, chronic leukaemia and acute leukaemia.
- 21. (Currently Amended) Use according to The method of claim 13 characterised characterized in that the disorder or disorders are selected from the group consisting of arthritis, restenosis; fibrotic disorders; mesangial cell proliferative disorders, diabetic nephropathy, malignant nephrosclerosis, thrombotic microangiopathy syndromes, organ transplant rejection, glomerulopathies,

metabolic disorders, inflammation and neurodegenerative diseases.

- 22. (Currently Amended) Use according to The method of claim 13, characterised characterized in that the disorder or disorders are selected from the group consisting of rheumatoid arthritis, inflammation, autoimmune disease, chronic obstructive pulmonary disease, asthma, inflammatory bowel disease, fibrosis, atherosclerosis, restenosis, vascular disease, cardiovascular disease, inflammation, renal disease and angiogenesis disorders.
- 23. (Currently Amended) Use A method of treatment comprising administering to a patient a the compound or compounds according to claim 1 as a kinase inhibitor.
- 24. (Currently Amended) Use according to The method of claim 23, characterised characterized in that the kinase is one or more raf-kinases, selected from the group consisting of A-Raf, B-Raf and Raf-1.
- 25. (Canceled)
- 26. (Canceled)
- 27. (Canceled)
- 28. (Currently Amended) Method for the treatment according to The method of claim 27 17, characterised characterized in that the disorder or disorders is cancerous cell growth mediated by one or more kinases.
- 29. (Currently Amended) Method A method for producing the compound or compounds of formula I claim 1, characterised in that comprising

a) a compound of formula II

$$(R^8)_p$$
 N
 N
 N
 R^7
 R^6

wherein

- L^1 is H or a metal ion, and R^6 , R^7 , R^8 and p are as defined in claim 1, is reacted
- b) with a compound of formula III,

$$L^{2} \xrightarrow{\qquad \qquad } X-Ar^{2}-(R^{10})_{r}$$

$$(R^{9})_{q}$$
III

wherein

L² is Cl, Br, $\frac{1}{I}$, OH, an esterified OH-group or a diazonium moiety, and Y, R^9 , q, X, Ar^2 , R^{10} and r are as defined in claim 1,

and optionally

c) isolating and/or treating the compound or compounds of claim 1 of formula I obtained by said reaction with an acid, to obtain the salt thereof.

30. (Currently Amended) Compound A compound or compounds of formula II,

$$(R^8)_p$$
 N
 N
 N
 R^7
 R^6

wherein

- L¹ is H or a metal ion, and R^6 , R^7 , R^8 and p are as defined in claim 1.
- 31. (Currently Amended) Compound A compound or compounds of formula III,

$$L^{2} \xrightarrow{\qquad \qquad } X-Ar^{2}-(R^{10})_{r} \qquad \qquad III$$

wherein

L² is Cl, Br, $\frac{1}{L}$, OH, an esterified OH-group or a diazonium moiety, and Y, R⁹, q, X, Ar², R¹⁰ and r are as defined in claim 1.